PTO/SB/08a Substitute for Form 1449A/PTO				Application Number	10/565,366
INFORMATION DISCLOSURE STATEMENT BY APPLICANT				Filing Date	January 23, 2006
				First Named Inventor	HARTUNG, Rolf
(Use as many sheets as necessary)				Art Unit	1764
				Examiner Name	To be assigned
Sheet	1	of	3	Attorney Docket	7601/84486

U.S. PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Document Number Number-Kind Code[2]	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear
/SY/	A1	US 4,788,322	11-29-1988	Merck & Co. Inc.	

FOREIGN PATENT DOCUMENTS

Examiner Initials*	Cite No.[1]	Foreign Patent Document Country Code[3]-Number[4]-Kind Code[5]	Publication Date MM-DD-YYYY	Name of Patentee or Applicant of Cited Document	Pages, Columns, Lines, Where Relevant Passages or Relevant Figures Appear	T[6]
/SY/	B1	EP 0 427 939	05-22-1991	American Cyanamid Co.		
/SY/	B2	EP 0 438 311	07-24-1991	Merck & Co. Inc.		
/SY/	B3	EP 0 823 416	08-07-1996	Ajinomoto Co., Inc.		
/SY/	B4	WO 91/07430	05-30-1991	The Upjohn Co.		

Examiner Signature		Date Considered	

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PTO/SB/08b Substitute for Form 1449B/PTO **INFORMATION DISCLOSURE STATEMENT BY APPLICANT** *(Use as many sheets as necessary)*				Application Number	10/565,366
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NON PATENT LITERATURE DOCUMENTS

Examiner Initials *	Cite No.[1]	Include name of the author (in CAPITAL LETTERS), title of the article (when appropriate), title of the item (book, magazine, journal, serial, symposium, catalog, etc.), date, page(s), volume-issue number(s), publisher, city and/or country where published.	T[2]
/SY/	C1	International Search Report for PCT/EP2004/006654 filed June 19, 2004.	
	C2	International Preliminary Report on Patentability for PCT/EP2004/006654 filed June 19, 2004.	
	C3	ALEXANDER, *et al.*, "A Diastereoselective Synthesis of (2S, 3R, 4S)-2-Amino-1-cyclohexyl-6-methylheptane-3,4-diol, The Abbott Aminodiol," *Tetrahedron Letters* 37:1961-1964 (1996).	
	C4	BIRCH, *et al.*, "The Structure and Some Reactions of the Iron Tricarbonyl Complex of Thebaine," *J. Chem. Soc. C.* 531 (1968).	
	C5	BLÁHA, *et al.*, "Stereoisomeric Chiral 2,9-Diazabicyclo[4.4.0.]Decane-3-,10-Diones as Models of Dipeptide Grouping: Synthesis, X-Ray, IR, NMR, and CD Studies," *Coll. Czech. Chem. Commun.* 49:712-742 (1984).	
	C6	CLINGMAN, *et al.*, "Effect of Amines on Hydrogenolysis of Alkylphenols," *J. Org. Chem.* 23:276-280 (February 1958).	
	C7	CORRINGER, *et al.*, "CCK-B Agonist of Antagonist Activities of Structurally Hindered and Peptidase-Resistant Boc-CCK_4 Derivatives," *J. Med. Chem.* 36:166-172 (1993).	
	C8	DEVANT, *et al.*, "Steroselektive Aldolreaktion Mit Chiralen Sekundären Acetamiden," *Chem. Ber.* 119:2191-2207 (1986).	
	C9	EISLER, *et al.*, "Amino Acids and Peptides. LXV. Analogues of Oxytocin," *Coll. Czech. Chem. Commun.* 31:4563-4580 (1966).	
	C10	FAUSTINI, et al., "Stereospecificity in the Transformation of α-Aminoacids into Fluroracids," *Tetrahedron Letters* 22:4533-4536 (1981).	
	C11	HAYASHI, *et al.*, "Chiral (β-Aminoalkyl)Phosphines. Highly Efficient Phosphine Ligands for Catalytic Asymmetric Grignard Cross-Coupling," *J. Org. Chem.* 48:2195-2202 (1983).	
	C12	HARRIS, *et al.*, "Structure of Ristocetin A: Configurational Studies of the Peptide," *J. Am. Chem. Soc.* 104: 363-365 (1982).	

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Sheet	3	of	3	Attorney Docket	7601/84486

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/SY/	C13	HOEKSTRA, *et al.*, "Large-Scale Synthesis of Anticoagulant Decapeptide MDL 28050," *Tetrahedron* 48:307-318 (1992).	
	C14	ISHIDA, *et al.*, "Micropeptins 88-A to 88-F, Chymotrypsin Inhibitors from the Cyanobacterium *Microcystis aeruginosa* (NIES-88)," *Tetrahedron* 54: 5545-5556 (1998).	
	C15	MINNAARD, *et al.*, "Synthesis of Enantiomerically Pure Cyclohexylglycine," *Synthetic Communications* 29(24): 4327-4332 (1999).	
	C16	PLATA, *et al.*, "The Stereospecific Preparation of an Hydroxyethylene Isotere Precursor via a Novel Piperidine-2,5-Dione Template," *Tetrahedron Letters* 32(30): 3623-3626 (1991).	
	C17	SCHUDA, *et al.*, "A Short and Efficient Synthesis of (3S, 4S)-4-[(*tert*-Butyloxycarbonyl)amino]-5-cyclohexyl-3-hydroxypentanoic Acid Ethyl Ester," *J. Org. Chem.* 53: 873-875 (1988).	
	C18	TAMURA, *et al.*, "Guanylpiperidine Peptidomimetics: Potent and Selective bis-Cation Inhibitors of Factor Xa," *Bioorg. Med. Chem. Lett.* 10(8): 745-749 (April 2000).	
	C19	TAMURA, *et al.*, "A Synthesis of Optically Active α-Cyclohexylglycine," *Synthetic Communications* 8(5): 345-351 (1978).	

Examiner Signature	/Shawquia Young/	Date Considered	08/17/2008

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